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Page 44, please delete all of Table 7a and insert therefor the attached  
substitute Table 7a.

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Page 45, please delete all of Table 7b and insert therefor the attached  
substitute Table 7b.

**IN THE CLAIMS:**

Please amend the claims as indicated below. A marked-up copy of the  
claims is attached.

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3. Nucleic acid according to Claim 1, which furthermore comprises a  
CDR2 region, selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

D I S Y S G S T K Y K P S L R S, (SEQ ID NO:35)

(b) a nucleotide sequence which encodes the amino acid sequence:

V I S Y D G S N K Y Y A D S V K G, (SEQ ID NO:36)

and

(c) a nucleotide sequence which encodes an amino acid sequence  
having an homology of at least 80% with an amino acid  
sequence from (a) or (b).

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6. Nucleic acid according to Claim 4, which furthermore comprises a CDR2 region selected from:

(a) a nucleotide sequence which encodes the amino acid sequence:

G S H Q R P S, (SEQ ID NO:41)

(b) a nucleotide sequence which encodes the amino acid sequence:

S N N Q R P S, (SEQ ID NO:42)

and

(c) a nucleotide sequence which encodes an amino acid sequence having an homology of at least 80% with an amino acid sequence from (a) or (b).

11. Vector, characterized in that it

(a) contains at least one copy of a nucleic acid according to Claim 1 and/or at least one copy of a nucleic acid according to Claim 4 or

(b) contains at least one copy of a nucleic acid according to Claim 7 and/or at least one copy of a nucleic acid according to Claim 9.

12. Cell, characterized in that it

(a) expresses a nucleic acid according to Claim 1 and/or a nucleic acid according to Claim 4 or

(b) a nucleic acid according to Claim 7 and/or a nucleic acid according to Claim 9.

*Sub C2*

13. Polypeptide, characterized in that it  
(a) is encoded by a nucleic acid according to Claim 1 and/or a nucleic acid according to Claim 4 or  
(b) by a nucleic acid according to Claim 7 and/or a nucleic acid according to Claim 9.

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16. Polypeptide according to Claim 13, characterized in that it is coupled to a labelling group or a toxin.

17. Antibody against a polypeptide according to Claim 13.

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*Sub C2*

19. Pharmaceutical composition which comprises, as the active component, a nucleic acid according to Claim 1, a vector according to Claim 11, a cell according to Claim 12, a polypeptide according to Claim 13 or an antibody according to Claim 17, where appropriate together with other active components and pharmaceutically customary adjuvants, additives or excipients.

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20. Use of a nucleic acid according to Claim 1, of a vector according to Claim 11, of a cell according to Claim 12, of a polypeptide according to Claim 13, of an antibody according to Claim 17, or of a pharmaceutical composition according to Claim 19 for preparing an agent for the diagnosis or for the treatment or prevention of AITP.